



CLAIMS

- Use of a multifunctional β-adrenergic receptor antagonist (β-blocker) compound comprising
 - i) a β-blocker component,
 - ii) at least one reactive oxygen species (ROS) scavenger component, and optionally
 - iii) at least one nitric oxide (NO) donor component in the preparation of a medicament.
- 2. Use of a multifunctional β -blocker compound according to claim 1, comprising
 - i) a β-blocker component,
 - ii) at least one ROS-scavenger component, and
 - iii) at least one nitric oxide (NO) donor component.
- Use according to claim 1, wherein said β-blocker component is selected from the group consisting of compounds used in medicine as β-adrenergic blockers, derivatives thereof, and compounds exhibiting affinity for βreceptors.
- 4. Use according to claim 1, wherein said ROS-scavenger component comprises an antioxidant reacting with ROS selected from the group consisting of superoxide, hydroxyl radicals, peroxynitrite, and hypochlorite.
- 5. Use according to claim 1, wherein said NO-donor comprises a group capable of providing nitric oxide in a form selected from uncharged and charged.
- 6. Use according to claim 4, wherein said ROS-scavenger component comprises a substituted N-oxide free radical.



20. Use according to any on of claims 15 to 18, wherein said multifunctional β -blocker compound has formula III

$$E \xrightarrow{D} \stackrel{H}{\underset{N}{\bigvee}} A$$

wherein A is C₁-C₄ alkyl or ROS-scavenger group;

B is selected from OH, O-NO2 and SH;

- D is H, or D is (CH₂)₂ and is connected to E and together with the neighboring atoms forms a 5-6 membered ring consisting of carbon atoms and one oxygen atom; and
- E is phenyl condensed with optionally substituted phenyl or optionally substituted 5-6 membered heterocycle containing one of -N-, -O-, and -S-S-; or
- E is thiadiazolyl substituted with morpholinyl or pyrrolidinyl-N-oxide, said morpholinyl being optionally substituted with one of OH, NO-donor group, and ROS-scavenger group, and said pyrrolidinyl-N-oxide group being bound to said thiadiazolyl vial -S- or via -CH₂-O-.
- 21. Use according to claim 23, wherein said compound is selected from the group consisting of compounds nos. 14, 15, 20-75, 1', 2', and 7'-24' as shown below.

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 - 22. A a multifunctional β-adrenergic receptor antagonist (β-blocker) compound comprising
 - a β-blocker component, i)
 - ii) at least one reactive oxygen species (ROS) scavenger component, and optionally
 - iii) at least one nitric oxide (NO) donor component for use as a medicament.
 - 23. A method of treating or preventing a disorder selected from the group consisting of disorders in which treatment with a β-antagonist is indicated, disorders associated with oxidative stress and free radical injury, and disorders in which treatment with a smooth muscle relaxant is indicated, in a mammal in need thereof, comprising administering to said mammal an effective amount of a multifunctional β-blocker compound comprising i) a βblocker component, ii) at least one reactive oxygen species (ROS) scavenger component, and optionally iii) at least one nitric oxide (NO) donor component.
 - 24. A method according to claim 23, wherein said disorder is selected from the group consisting of ischemia, ischemia-reperfusion tissue injury, acute and chronic inflammatory conditions, angina, atherosclerosis, impotence, hypertension, pulmonary hypertension, systemic hypertension, obesity or pregnancy-induced hypertension, palpitations, arrhythmias, cardiomyopathy, congestive heart failure, hyperthyroidism, anxiety, tremor, migraine, alcohol withdrawal, tachycardia, thyrotoxicosis, pheochromocytoma, esophageal varices, glaucoma, conditions associated with excess intraocular fluid, diabetes mellitus, and carcinogenesis.
 - 25. A method according to claim 23, wherein said administration or treatment is selected from the group consisting of topical, oral, and parenteral.





- 26. A method according to claim 23, wherein said administration or treatment is selected from the group consisting of suppository, by way of injection, and by way of infusion.
- 27. A method according to claim 23, wherein said multifunctional β-blocker compound is administered by a route selected from intramuscular, intraperitoneal, intravenous, ICV, intracisternal injection or infusion, subcutaneous injection, implant, inhalation spray, nasal, vaginal, rectal, sublingual, and urethral.
- 28. A method according to claim 23, wherein said mammal is human.
- 29. A multifunctional β-adrenergic receptor antagonist compound comprising
 - i) a β-blocker component,
 - at least one ROS-scavenger component, and optionally
 - iii) at least one NO-donor component.
- 30. A multifunctional antagonist according to claim 29, wherein said β-blocker component is selected from the group consisting of compounds used in medicine as β-adrenergic blockers, derivatives thereof, and compounds exhibiting affinity for β-receptors.
- 31. A multifunctional antagonist according to claim 29, wherein said ROS-scavenger component comprises an antioxidant reacting with ROS selected from the group consisting of superoxide, hydroxyl radicals, peroxynitrite, and hypochlorite.
- 32. A multifunctional antagonist according to claim 29, wherein said ROS-scavenger component comprises any of alkenyl group, aryl group, substituted aryl group, sulfhydryl, dithiol in oxidized or reduced form, and group that is converted *in vivo* into a sulfhydryl in its oxidized or reduced form.



- i) a dosage amount of at least one compound having β -blocker activity and ROS-scavenging activity,
- ii) instructions for use; and
- iii) optionally means for the delivery of said compound.